

## COMBINATIONS FOR THE TREATMENT OF DISEASES INVOLVING ANGIOGENESIS

This invention relates to a method for treating diseases involving active angiogenesis,  
5 to compositions useful for the treatment of diseases involving angiogenesis and to the  
use of the compositions in the preparation of a medicament for the treatment of  
diseases involving active angiogenesis. In one aspect of the invention the method  
involves the administration to a mammal of an inhibitor of nitric oxide in combination  
with a compound inducing vascular damage.

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Formation of new vasculature by angiogenesis is a key pathological feature of several  
diseases (J Folkman, New England Journal of Medicine 333, 1757-1763, 1995). For  
example, for a solid tumour to grow it must develop its own blood supply upon which  
it depends critically for the provision of oxygen and nutrients; if this blood supply is  
15 mechanically shut off the tumour undergoes necrotic death. Neovascularisation is also  
a clinical feature of skin lesions in psoriasis, of the invasive pannus in the joints of  
rheumatoid arthritis patients and of atherosclerotic plaques. Retinal neovascularisation  
is pathological in macular degeneration and in diabetic retinopathy. In all these diseases  
reversal of neovascularisation by damaging the newly-formed vascular endothelium is  
20 expected to have a beneficial therapeutic effect.

Certain chemical compounds have been shown to have vascular damaging activity  
against the newly formed endothelium of solid tumours. These agents include, for  
example, combretastatin A4 phosphate (Dark et al., Cancer Research 57, 1829-1834,  
25 1997), combretastatin analogues (for example those described in J Med Chem 41,  
3022-32, 1998 by Ohsumi et al.), the flavone acetic acids, for example 5,6-  
dimethylxanthenone acetic acid (Zwi, Pathology, 26, 161-9, 1994), colchicine  
(Baguley et al. Eur J Cancer 27, 482-7, 1991). However some tumours are resistant to  
these agents.

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One characteristic of tumours relatively resistant to vascular damaging agents is their  
ability to produce large amounts of nitric oxide. The role of nitric oxide in tumour

growth is unclear and there have been reports of both tumour-stimulating and tumour-inhibiting effects (Chinje and Stratford, Essays Biochem. 32, 61-72, 1997). It has been suggested that the antitumour effects of 5,6-dimethylxanthenone acetic acid are mediated in part by nitric oxide production (Thompson et al. Cancer Chemother  
5 Pharmacol. 31, 151-5, 1992).

WO-A 9509621 and Br. J Cancer (1998), 77(3), 426-433 disclose combinations of cytokine releasing anticancer agents (TNF-releasing agents). These relate to ameliorating the effects of pro-inflammatory cytokines. There is no suggestion of  
10 synergistic activity from a combination of a vascular damaging agent (many of which have no pro-inflammatory activity) and an NO inhibitor.

⇒ We have found that the efficacy of vascular damaging agents can be improved by combining the treatment with inhibitors of the formation or action of nitric oxide in a  
15 mammalian system.

In particular the efficacy of vascular damaging agents can be improved by combination with inhibitors of nitric oxide synthases, the enzymes that produce nitric oxide from arginine. In particular the efficacy of vascular damaging agents against tumours  
20 relatively resistant to their effects is improved by treatment with a nitric oxide synthase inhibitor.

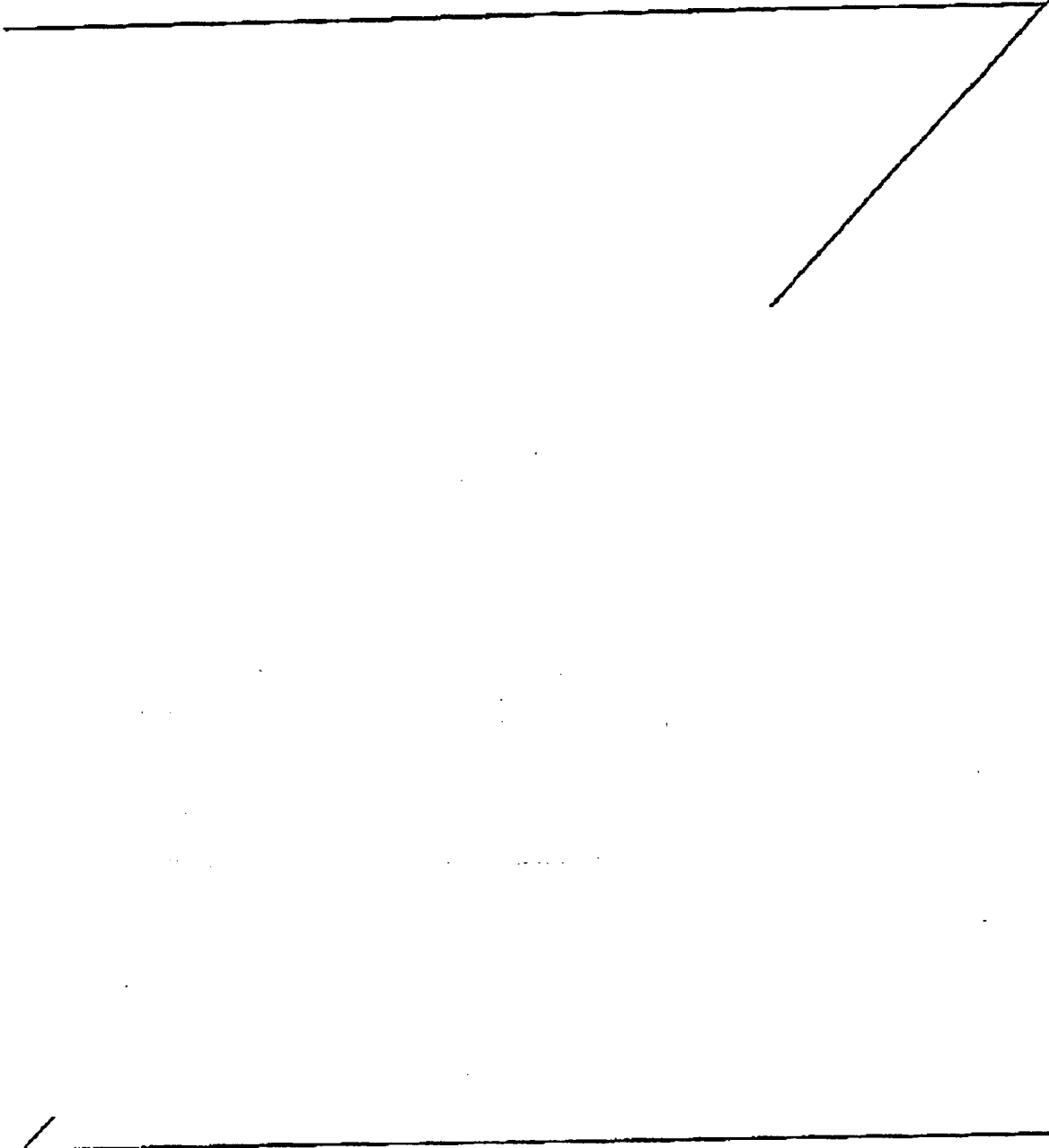
Accordingly in one aspect of the invention we provide a method of treatment for a mammal having a disease that involves active angiogenesis such method comprising the  
25 administration of a therapeutic or subtherapeutic amount of a vascular damaging agent together with an inhibitor of nitric oxide synthase in an amount sufficient to augment the effect of the vascular damaging agent. The method is useful for the treatment of diseases such as cancers, especially solid tumours, psoriasis, diabetic retinopathy, macular degeneration, atherosclerosis and rheumatoid arthritis.

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The vascular damaging agent and the nitric oxide synthase inhibitor can be administered together or separately. The method may be used as a sole therapy or in

combination with other treatments. For the treatment of solid tumours compounds of the invention may be administered in combination with radiotherapy or in combination with other anti-tumour substances for example those selected from mitotic inhibitors, for example vinblastine, paclitaxel and docetaxel; alkylating agents, for example

5 cisplatin, carboplatin and cyclophosphamide; antimetabolites, for example 5-fluorouracil, cytosine arabinoside and hydroxyurea; intercalating agents for example



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adriamycin and bleomycin; enzymes, for example asparaginase; topoisomerase inhibitors for example etoposide, topotecan and irinotecan; thymidylate synthase inhibitors for example raltitrexed; biological response modifiers for example interferon; antibodies for example edrecolomab; and anti-hormones for example tamoxifen. Such  
5 combination treatment may involve simultaneous or sequential application of the individual components of the treatment.

The vascular damaging agent and the nitric oxide synthase inhibitor can be administered by the same route or by different routes. Such routes of administration  
10 include oral, buccal, nasal, topical, rectal and parenteral administration. Each component of the method, the vascular damaging agent and the nitric oxide synthase inhibitor may independently be administered in a form suitable for the intended route of administration and such forms may be prepared in a conventional manner using conventional excipients. For example for oral administration the pharmaceutical  
15 compositions may take the form of tablets or capsules. For nasal administration or administration by inhalation the compounds may be conveniently delivered as a powder or in solution. Topical administration may be as an ointment or cream and rectal administration may be as a suppository. For parenteral injection (including intravenous, subcutaneous, intramuscular, intravascular or infusion) the composition may take the  
20 form of, for example, a sterile solution, suspension or emulsion.

The preferred route of administration of each component will depend on the disease being treated. For solid tumours the components may each advantageously be delivered, either together or separately, as an intravenous infusion.

25 Vascular damaging agents are compounds which induce selective damage to newly formed, rather than established, vasculature. Many such compounds are known and it is considered this invention is generally applicable to such agents. Such agents include tubulin-binding agents, for example the combretastatins and their prodrugs, the colchicins and their prodrugs and (Z)-2-methoxy-5-[2-(3,4,5-  
30 trimethoxyphenyl)vinyl]phenylamine and its prodrugs, TNF-alpha inducing agents such as the xanthenone acetic acids, for example dimethylxanthenoneacetic acid, and antibodies targeted to the vasculature.

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A wide variety of compounds which inhibit the formation or action of nitric oxide in mammalian systems can be employed. Specifically nitric oxide synthase inhibitors are those compounds which inhibit any of the forms of nitric oxide synthase. Such agents include derivatives of arginine, ornithine, lysine and citrulline, S-alkylthioureas and aminoguanidines. Where the nitric oxide synthase inhibitor is a derivative of arginine it may be, for example, an N<sup>G</sup>-substituted L-arginine selected from N<sup>G</sup>-nitro-L-arginine and alkyl esters thereof, N<sup>G</sup>-methyl-L-arginine and N<sup>G</sup>-amino-L-arginine. Where the nitric oxide synthase inhibitor is a derivative of ornithine it may be, for example L-N6-(1-iminoethyl)-ornithine. Where the nitric oxide synthase inhibitor is a derivative of lysine it may be, for example L-N6-(1-iminoethyl)-lysine. Where the nitric oxide synthase inhibitor is a derivative of citrulline it may be, for example L-thiocitrulline, L-homothiocitrulline or an S-alkylthiocitrulline such as S-methyl-L-thiocitrulline.

In a further embodiment of the invention there is provided a composition for the treatment of diseases involving active angiogenesis. The composition of the invention comprises a vascular damaging agent in combination with a nitric oxide synthase inhibitor where both the vascular damaging agent and the nitric oxide synthase inhibitor are as hereinbefore defined.

Thus for example the composition may contain for example a combretastatin derivative, a colchicine derivative, a colchinel derivative, a xanthenone acetic acid derivative or a vascular targeted antibody, in combination with a nitric oxide synthase inhibitor for example a derivative of arginine, a derivative of ornithine, a derivative of lysine, a derivative of citrulline, a S-alkylthioureas or an aminoguanidine.

Particular examples of vascular damaging agents that may be present in the composition include combretastatin A4 and its prodrugs for example combretastatin A4 phosphate, (Z)-2-methoxy-5-[2-(3,4,5-trimethoxyphenyl)vinyl]phenylamine and its prodrugs, N-acetylcolchinel and its prodrugs for example N-acetylcolchinel-O-phosphate and 5,6-dimethylxanthenoneacetic acid.

Particular examples of nitric oxide synthase inhibitors which may be present in the composition include derivatives of arginine, ornithine, lysine and citrulline, S-alkylthioureas aminoguanidines and aminopyridines. Where the nitric oxide synthase inhibitor is a derivative of arginine it may be, for example, an N<sup>G</sup>-substituted L-arginine selected from N<sup>G</sup>-nitro-L-arginine and alkyl esters thereof, N<sup>G</sup>-methyl-L-arginine and N<sup>G</sup>-amino-L-arginine. Where the nitric oxide synthase inhibitor is a derivative of ornithine it may be, for example L-N6-(1-iminoethyl)-ornithine. Where the nitric oxide synthase inhibitor is a derivative of lysine it may be, for example L-N6-(1-iminoethyl)-lysine. Where the nitric oxide synthase inhibitor is a derivative of citrulline it may be, for example L-thiocitrulline, L-homothiocitrulline or an S-alkylthiocitrulline such as S-methyl-L-thiocitrulline. Where the nitric oxide synthase inhibitor is an aminopyridine it may be for example 2-amino-4-methylpyridine.

The composition is useful for the treatment of diseases involving active angiogenesis for example solid tumours, psoriasis, diabetic retinopathy, macular degeneration, atherosclerosis and rheumatoid arthritis.

The relative proportion of each component will be determined by the identity of each individual vascular damaging agent or nitric oxide synthase inhibitor and by the disease to be treated.

The composition may include pharmaceutically acceptable excipients selected with regard to the intended route of administration and standard pharmaceutical practice. The composition may take a form suitable for oral, buccal, nasal, topical, rectal or parenteral administration and may be prepared in a conventional manner using conventional excipients. For example for oral administration the composition may take the form of tablets or capsules. For nasal administration or administration by inhalation the compounds may be conveniently delivered as a powder or in solution. Topical administration may be as an ointment or cream and rectal administration may be as a suppository. For parenteral injection (including intravenous, subcutaneous, intramuscular, intravascular or infusion) the composition may take the form of, for example, a sterile solution, suspension or emulsion.

- The dose of a compound of the invention required for the prophylaxis or treatment of a particular condition will vary depending on the identity of the individual components, the route of administration, the form and severity of the condition and whether the compound is to be administered alone or in combination with another drug. Thus the precise dose will be determined by the administering physician and will depend on the particular vascular damaging agent and NO synthase inhibitor in the composition. However the dose of the vascular damaging agent envisaged is, for example, in the range 10-1000mg/m<sup>2</sup> body surface, preferably 20-200mg/ m<sup>2</sup> and that for the nitric oxide inhibitor 1-1000mg/m<sup>2</sup>, preferably 5-500mg/ m<sup>2</sup>. A unit dose form of the vascular damaging agent as, for example, a sterile solution for injection will usually contain, for example, 40-400mg of the active ingredient. A unit dose form of the nitric oxide synthase inhibitor as, for example, a sterile solution for injection will usually contain, for example, 10-1000mg of the active ingredient. A unit dose form of a composition containing both a vascular damaging agent and a nitric oxide synthase inhibitor as, for example, a sterile solution for injection will usually contain, for example, 40-400mg of the vascular damaging agent and 10-1000mg of the nitric oxide synthase inhibitor.
- The composition of the invention may be administered as a sole therapy or in combination with other treatments. For the treatment of solid tumours the composition may be administered in combination with radiotherapy or in combination with other anti-tumour substances for example those selected from mitotic inhibitors, for example vinblastine, paclitaxel and docetaxel; alkylating agents, for example cisplatin, carboplatin and cyclophosphamide; antimetabolites, for example 5-fluorouracil, cytosine arabinoside and hydroxyurea; intercalating agents for example adriamycin and bleomycin; enzymes, for example asparaginase; topoisomerase inhibitors for example etoposide, topotecan and irinotecan; thymidylate synthase inhibitors for example raltitrexed; biological response modifiers for example interferon; antibodies for example edrecolomab; and anti-hormones for example tamoxifen. Such combination treatment may involve simultaneous or sequential application of the individual components of the treatment.

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In a further embodiment of the invention we provide the use of a composition of the invention for the preparation of a medicament for the treatment of a disease involving active angiogenesis.

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The invention will now be illustrated by the following Examples in which biological assays are used to illustrate the invention:

10 Induction of necrosis

Mice bearing either CaNT or SaS tumours were treated with the test compound and tumours excised after 24h, fixed in formalin, embedded in paraffin, sectioned and stained with haematoxylin and eosin. Sections were scored based on area of necrosis as follows:

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% necrosis	score	% necrosis	score
0-10	1	51-60	6
11-20	2	61-70	7
21-30	3	71-80	8
31-40	4	81-90	9
41-50	5	91-100	10

Control tumours had mean scores of 2.0 (CaNT) and 1.0 (SaS).

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EXAMPLE 1

In this assay the effect of a given dose of either a vascular damaging agent or a nitric oxide synthase inhibitor administered alone can be compared with the effect of a combination of the two agents.

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**Table 2: Enhancement of Combretastatin A4 phosphate (CA4P) activity in SaS tumours by coadministration of 2-amino-4-methylpyridine (AMP)**

Treatment	Necrosis score ±SEM (n)
None	1.0±0 (10)
CA4P, 500mg/kg	1.7±0.7 (3)
AMP, 10mg/kg	1.0 (2)
CA4P, 500mg/kg + AMP 10mg/kg	4.5 (2)

## EXAMPLE 3

Activity against tumour vasculature measured by fluorescent dye.

- The following experiment further demonstrates the ability of the compounds to
- 5 damage tumour vasculature.
- Tumour functional vascular volume in CaNT tumour-bearing mice was measured using the fluorescent dye Hoechst 33342 according to the method of Smith *et al* (Brit J Cancer 57, 247-253, 1988). The fluorescent dye was dissolved in saline at 6.25 mg/ml and injected intravenously at 10 mg/kg 24 hours after intra peritoneal drug treatment.
- 10 One minute later, animals were killed and tumours excised and frozen; 10  $\mu$ m sections were cut at 3 different levels and observed under UV illumination using an Olympus microscope equipped with epifluorescence. Blood vessels were identified by their fluorescent outlines and vascular volume was quantified using a point scoring system based on that described by Chalkley, (J Natl Cancer Inst, 4, 47-53, 1943). All
- 15 estimates were based on counting a minimum of 100 fields from sections cut at the 3 different levels.

Table 3: Enhancement of Combretastatin A4 phosphate (CA4P) activity in CaNT tumours by coadministration of L-N<sup>G</sup>-nitroarginine (L-NNA).

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Treatment	Vascular Volume % $\pm$ SEM (n)
None	2.35
CA4P, 25mg/kg	1.03 $\pm$ 0.14 (4)
L-NNA, 10mg/kg	2.45 $\pm$ 0.04 (3)
CA4P, 25mg/kg + L-NNA 10mg/kg	0.63 $\pm$ 0.25 (3)